FORM 6-K

SECURITIES AND EXCHANGE COMMISSION Washington, D.C. 20549

Report of Foreign Private Issuer

Pursuant to Rule 13a-16 or 15d-16 under the Securities Exchange Act of 1934

| For the month of September | 2003 |
|----------------------------|---------|
| Commission File Number | 0-16174 |

TEVA PHARMACEUTICAL INDUSTRIES LIMITED

(Translation of registrant's name into English)

5 Basel Street, P.O. Box 3190 Petach Tikva 49131 Israel (Address of principal executive offices)

| Indicate by check mark whether the registrant files or v Form 40-F: | vill file annual reports under cover of Form 20-F or | |
|---|--|--|
| Form 20-F <u>X</u> | Form 40-F | |
| Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1): | | |
| Indicate by check mark if the registrant is submitting the Rule 101(b)(7): | ne Form 6-K in paper as permitted by Regulation S-T | |
| Indicate by check mark whether by furnishing the information contained in this Form, the registrant is also hereby furnishing the information to the Commission pursuant to Rule 12g3-2(b) under the Securities Exchange Act of 1934. | | |
| Yes | No <u>X</u> | |
| If "Yes" is marked, indicate below the file number assisting 12g(3)-2(b): 82 | gned to the registrant in connection with Rule | |



Contact: Dan Suesskind

Chief Financial Officer

Teva Pharmaceutical Industries Ltd

(011) 972-2-589-2840

Bill Fletcher

President and CEO Teva North America. (215) 591-8800

FOR IMMEDIATE RELEASE

Dorit Meltzer

Director, Investor Relations Teva Pharmaceutical Industries Ltd. (011) 972-3-926-7554

TEVA ANNOUNCES APPROVAL OF NEFAZODONE HCI TABLETS 50 mg, 100 mg, 150 mg, 200 mg and 250 mg

Jerusalem, Israel, September 16, 2003 – Teva Pharmaceutical Industries Ltd. (Nasdaq: TEVA) announced today that the U.S. Food and Drug Administration has granted approval of the company's ANDA for Nefazodone Hydrochloride Tablets, 50 mg, 100 mg, 150 mg, 200 mg and 250 mg. Shipment of this product will begin immediately.

Nefazodone HCl Tablets are the AB-rated generic equivalent of Bristol-Myers Squibb's Serzone[®] Tablets indicated for the treatment of depression.

The brand product has annual sales of approximately \$220 million.

Teva Pharmaceutical Industries Ltd, headquartered in Israel, is among the top 30 pharmaceutical companies in the world. The company develops, manufactures, and markets generic and branded human pharmaceuticals and active pharmaceutical ingredients. Close to 90 percent of Teva's sales are in North America and Europe.

Safe Harbor Statement under the U. S. Private Securities Litigation Reform Act of 1995: This release contains forward-looking statements, which express the current beliefs and expectations of management. Such statements are based on current expectations and involve a number of known and unknown risks and uncertainties that could cause Teva's future results, performance or achievements to differ significantly from the results, performance or achievements expressed or implied by such forward-looking statements. Important factors that could cause or contribute to such differences include Teva's ability to successfully develop and commercialize additional pharmaceutical products, the introduction of competitive generic products, the impact of competition from brand-name companies that sell their own generic products or successfully extend the exclusivity period of their branded products, Teva's ability to rapidly integrate the operations of acquired businesses, the availability of product liability coverage in the current insurance market, the impact of pharmaceutical industry regulation and pending legislation that could affect the pharmaceutical industry, the difficulty of predicting U.S. Food and Drug Administration ("FDA") and other regulatory authority approvals, the regulatory environment and changes in the health policies and structure of various countries, acceptance and demand for new pharmaceutical products and new therapies, uncertainties regarding market acceptance of innovative products newly launched, currently being sold or in development, the impact of restructuring of clients, reliance on strategic alliances, exposure to product liability claims, dependence on patent and other protections for innovative products, fluctuations in currency, exchange and interest rates, operating results and other factors that are discussed in Teva's Annual Report on Form 20-F and its other filings with the U.S. Securities and Exchange Commission ("SEC"). Forward-looking statements speak only as of the date on which t

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

TEVA PHARMACEUTICAL INDUSTRIES LIMITED (Registrant)

/s/ Dan Suesskind Name: Dan Suesskind Title: Chief Financial Officer

Date: September 16, 2003